This listing of claims will replace all prior versions and listings of claims in this

application:

Listing of Claims

Claims 1–36 (canceled)

Claim 37 (currently amended): A method of processing sulfide minerals and concentrates

by oxidation of sulfide minerals in an aqueous medium using an oxidizing agent which is

one or more of nitric acid, nitrous acid and their oxides, the method comprising:

subjecting in an oxidation reactor a slurry containing the sulfide minerals to

oxidation under agitation and under controlled conditions of slurry acidity using the

oxidizing agent which is one or more of nitric acid, nitrous acid and their oxides;

forming in the oxidation reactor a sulfuric acid as a result of the sulfide oxidation;

constantly neutralizing the sulfuric acid in the oxidation reactor using an acidity

neutralizer to an acidity level at which no formation of elementary sulfur occurs;

removing of heat released during the sulfide oxidation from the oxidation reactor;

transferring NO from the oxidation reactor into a regeneration oxidizer;

regenerating N₂O₃ from the transferred NO using air or oxygen in the

regeneration oxidizer; and

transferring the regenerated N₂O₃ into the oxidation reactor;

wherein the temperature in the oxidation reactor is maintained in a range from 20

to 90 °C and

wherein a liquid-to-solid ratio in the slurry in the oxidation reactor is between 1:1

to 5:1.

Claim 38 (previously presented): The method according to claim 37 in which the acidity

neutralizer is one or more of CaCO₃, MgCO₃, Ca(OH)₂, CaO, NaOH and CaHPO₄.

Claim 39 (currently amended): The method according to claim 37 in which the

temperature in the oxidation reactor is maintained in the range of 65-85°C.

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Claim 40 (currently amended): The method according to claim 37, further comprising

before transferring the regenerated N₂O₃ into the oxidation reactor, separating the

N₂O₃, formed in said method, from N₂ by absorbing the N₂O₃ from a mix of gases

comprising N₂ and N₂O₃ into a sulfuric acid solution which has a concentration in the

range 75-98%; and

denitrating releasing N₂O₃ from the sulfuric acid solution thermally by heating it

to a temperature not exceeding 250°C, and/or chemically by introduction of a denitrating

substance.

Claim 41 (previously presented): The method according to claim 40, in which the

denitrating substance is one or more of an alcohol, formaldehyde and other chemical

reducing agents.

Claim 42 (previously presented): The method according to claim 37, further comprising

separating the NO, formed in said method, from N₂ by absorbing the NO from a

mix of gases comprising N₂ and NO into a monovalent copper salt solution;

denitrating the monovalent copper salt solution using a dosed supply of

compressed air, with optional simultaneous heating of the solution.

Claim 43 (previously presented): The method according to claim 42 in which the

monovalent copper salt solution contains a stabilizing agent to impede oxidation of

copper from monovalent to bivalent.

Claim 44 (previously presented): The method according to claim 43 in which the

stabilizing agent is one or more of tributyl phosphate, adipodinitrile, or reducing agents

such as formaldehyde or hydrazine.

Claim 45 (canceled)

Claim 46 (previously presented): The method according to claim 37, wherein the

regenerating the N₂O₃ from the NO formed in said method is performed using pure

oxygen in an individual regeneration oxidizer and at a temperature of 15-25°C.

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